ANTIBACTERIAL FOAMING CRANBERRY- triclos an liquid MEIJER DISTRIBUTION INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

ACTIVE INGREDIENT

TRICLOSAN 0.46%

PURPOSE

ANTIBACTERIAL

USES

HELPS REDUCE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER.

STOP USING THIS PRODUCT AND ASK DOCTOR

IF IRRITATION AND REDNESS DEVELOP AND LAST

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

PUMP ONTO DRY HANDS, WORK INTO A LATHER AND RINSE THOROUGHLY.

OTHER INFORMATION

STORE AT ROOM TEMPERATURE

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM XYLENE SULFONATE, DIPROPYLENE GLYCOL, GLYCERIN, SODIUM PCA, AMMONIUM LAURYL SULFATE, COCAMIDOPROPYL BETAINE, POLYQUATERNIUM-10, FRAGRANCE (PARFUM), VACCINIUM MACROCARPON (CRANBERRY) FRUIT EXTRACT, TOCOPHERYL ACETATE, CAMELLIA OLEIFERA (TEA) LEAF EXTRACT, ALOE BARBADENSIS (ALOE VERA) LEAF EXTRACT, DISODIUM PHOSPHATE, CETYL ALCOHOL, PROPYLENE GLYCOL, CITRIC ACID, METHYLPARABEN,

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ANTIBACTERIAL FOAMING CRANBERRY

triclosan liquid

Product Information	Prod	luct	Information
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-186

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.46 mL in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
SODIUM XYLENESULFONATE (UNII: G4LZF950 UR)				
DIPRO PYLENE GLYCOL (UNII: E107L85C40)				
GLYCERIN (UNII: PDC6A3C0OX)				
SODIUM PYRROLIDONE CARBO XYLATE (UNII: 469 OTG57A2)				
AMMO NIUM LAURYL SULFATE (UNII: O7AO2R1M0B)				

CO CAMIDO PRO PYL BETAINE (UNII: 50 CF3011KX)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
CRANBERRY (UNII: 0MVO31Q3QS)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
SODIUM PHO SPHATE (UNII: SE337SVY37)	
C12-16 ALCOHOLS (UNII: S4827SZE3L)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:41250-186-08	221 mL in 1 BOTTLE, PUMP			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	11/18/2011		

Labeler - MELIER DISTRIBUTION INC (006959555)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture	

Revised: 11/2011 MEIJER DISTRIBUTION INC